Original Paper

COVID-19–Relevant Insights Into the Elevated Risk of Accidental Injuries in Survivors of SARS and Their Relatives in Taiwan: Retrospective Cohort Study

Chieh Sung^{1,2*}, PhD; Chi-Hsiang Chung³, PhD; Chien-An Sun^{4,5}, PhD; Chang-Huei Tsao^{2,6}, PhD; Daphne Yih Ng⁷, MS; Tsu-Hsuan Weng^{2,8}, MS; Li-Yun Fann^{9,10}, PhD; Fu-Huang Lin³, PhD; Wu-Chien Chien^{2,3,8*}, PhD

Corresponding Author:

Li-Yun Fann, PhD
Department of Nursing
Taipei City Hospital
145 Zhengzhou Road, Datong District
Taipei, 103212
Taiwan

Phone: 886 0225553000 Email: fanly99@gmail.com

Abstract

Background: The 2003 outbreak of severe acute respiratory syndrome (SARS), caused by a novel coronavirus, heavily impacted Taiwan's health care system, triggering clinical crises and lasting effects among affected individuals and families. The first case in Taiwan was identified on February 25, 2003, and the final case was reported on June 15, 2003. During the epidemic, 346 people were diagnosed with SARS, leading to 37 deaths. Outbreaks also occurred in China, Singapore, and Toronto (Canada), showing the vulnerability of global health systems to new zoonotic diseases. Clinically, SARS causes high fever and severe lung inflammation. Survivors often had long-term lung problems, including fibrosis, and bone issues like osteonecrosis, mostly due to high-dose steroid treatment. Although studies have looked at long-term outcomes—especially lung and bone issues—none followed patients beyond 7 years. The COVID-19 pandemic further revealed gaps in understanding how serious viral infections affect wider health areas, including unintentional and intentional injuries. Data on related hospitalizations also remain limited.

Objective: This study aimed to investigate the long-term risk of both unintentional and intentional injuries among survivors of SARS and their relatives, using a nationwide population-based cohort.

Methods: This retrospective cohort study used data from Taiwan's National Health Insurance Research Database, focused on 285 individuals diagnosed with SARS in 2003 and 699 of their relatives, matched in a 1:10 ratio with controls. Injury risks were assessed using Fine and Gray's competing risk models, adjusting for sociodemographic and clinical covariates, over a follow-up period of up to 15 years.

Results: Survivors of SARS exhibited a significant increase in the risk of accidental injuries, with an adjusted hazard ratio (AHR) of 1.631 (95% CI 1.184-2.011; *P*<.001), indicating persistent physiological vulnerabilities postinfection. Family members of survivors of SARS also had elevated injury risk (AHR 1.572, 95% CI 1.148-1.927; *P*<.001), possibly due to stress and caregiving burdens. Subgroup analysis showed increased risks for poisoning (AHR 2.701, 95% CI 1.956-4.084;

¹Department of Senior Citizen Care and Welfare, Deh Yu College of Nursing and Health, Keelung City, Taiwan

²Department of Medical Research, Tri-Service General Hospital, National Defense Medical Center, Taipei, Taiwan

³School of Public Health, National Defense Medical Center, Taipei, Taiwan

⁴Department of Public Health, College of Medicine, Fu-Jen Catholic University, New Taipei City, Taiwan

⁵Big Data Research Center, College of Medicine, Fu-Jen Catholic University, New Taipei City, Taiwan

⁶Department of Microbiology & Immunology, National Defense Medical Center, National Defense Medical Center, Taipei, Taiwan

⁷Department of Family Medicine, Tri-Service General Hospital, National Defense Medical Center, Taipei, Taiwan

⁸Graduate Institute of Medical Sciences, National Defense Medical Center, Taipei, Taiwan

⁹Department of Nursing, Taipei City Hospital, Taipei, Taiwan

¹⁰Department of Nursing, Tzu Chi University, Hualien, Taiwan

^{*}these authors contributed equally

P<.001) and falls (AHR 1.524, 95% CI 1.102-1.878; P=.003) among survivors. Relatives faced higher risks for traffic incidents (AHR 2.003, 95% CI 1.462-2.459), poisoning (AHR 1.531, 95% CI 1.120-1.886), medical incidents, falls (AHR 1.802, 95% CI 1.324-2.214), and crushing injuries (AHR 2.469, 95% CI 1.803-3.026; all P<.001). These findings highlight the need for targeted preventive measures to address long-term health risks in both survivors of SARS and their families.

Conclusions: Survivors of SARS and their relatives face increased injury risks, highlighting long-term physical and psychosocial vulnerabilities after severe infectious outbreaks. These findings suggest that health care systems should provide preventive and supportive measures to mitigate long-term impacts for those affected by pandemics.

JMIR Public Health Surveill 2025;11:e70608; doi: 10.2196/70608

Keywords: SARS; accidental injuries; COVID-19 pandemic; psychosocial impacts; intentional injuries; survivors of SARS; accidental falls; poisoning; psychological stress; caregivers; population surveillance; severe acute respiratory syndrome

Introduction

An outbreak of severe acute respiratory syndrome (SARS) caused by a novel coronavirus severely affected Taiwan in 2003 [1], with the first confirmed diagnosis of SARS on February 25, 2003, and the last diagnosis on June 15, 2003. Through this endemic outbreak, there were 346 patients diagnosed with SARS, and 37 died among these patients [2], and there were also SARS outbreaks in China, Singapore, and Toronto, Canada, at about the same time [2]. The COVID-19 pandemic has had a global impact on all facets of medicine, resulting in significant stress and challenges in the medical community.

The main clinical manifestations after infection were high fever and severe lung inflammation [3,4]. The patients who survived had residual pulmonary fibrosis, as well as osteonecrosis, resulting from treatment with large doses of steroid pulse therapy [5]. Several studies have investigated the long-term outcomes of recovered patients with SARS, especially with respect to lung and bone damage, but no follow-up periods of more than 7 years have been reported [6,7]. Furthermore, there is a scarcity of research regarding the effects of COVID-19 on unintentional injuries and intentional injuries, as well as admission rates [8]. Survivors of respiratory infections like SARS and COVID-19 frequently report persistent symptoms that severely impact their daily functioning and quality of life. These symptoms include chronic fatigue, muscle weakness, and neurological manifestations such as "brain fog" and cognitive deficits, which are well-documented consequences of both diseases [9-11]. Beyond the physical impairments, there is growing evidence highlighting the long-term mental health burden of these infections, with elevated rates of anxiety, depression, posttraumatic stress disorder, and sleep disturbances reported to persist for months or even years after recovery [12-14]. These multifaceted impairments may heighten the risk of accidental injuries in affected individuals. Specifically, cognitive dysfunction, such as reduced attention, slower processing speed, and impaired judgment, is known to increase vulnerability to various types of accidents, including falls and motor vehicle incidents [15,16]. Moreover, physical limitations like chronic fatigue and muscle weakness directly compromise balance, coordination, and reaction time, thus elevating the likelihood of falls, particularly in older adults [17,18]. The increased prevalence of mental health disorders postpandemic may also lead to greater reliance on psychotropic medications, which carry their own risks, including sedation and impaired gait, further contributing to the elevated risk of falls [19]. While the direct link between SARS and subsequent accidental injuries has been underexplored, the parallels in the long-term physical and psychological consequences observed in survivors of COVID-19 strongly suggest that survivors of SARS may similarly face increased vulnerability to injuries [20,21].

The implementation of social distancing and quarantine measures during COVID-19 likely led to a decrease in trauma surgery cases, as restrictions reduced high-risk activities [8]. We hypothesized that SARS is associated with an increased risk of these injuries and tested this hypothesis through a nationwide cohort study using the National Health Insurance Research Database (NHIRD), which includes claims data for Taiwan's entire population [22].

Methods

Data Sources

Established in 1995, Taiwan's National Health Insurance (NHI) Program is a mandatory and universal health insurance system that covers approximately 23 million beneficiaries—over 99% of the population [22]. The NHIRD contains comprehensive and detailed data regarding the total outpatients and inpatients [23-33].

We selected an inpatient dataset from 2000 to 2015 from the NHIRD, with individual diagnoses coded using the *International Classification of Diseases, Ninth Revision, Clinical Modification (ICD-9-CM)*. The details of the NHI program have been documented in several previous studies [34-39]. Several studies have also confirmed the precision and validity of these diagnoses [40-42].

Ethical Considerations

This study was conducted in accordance with the Code of Ethics of the World Medical Association (Declaration of Helsinki). The Institutional Review Board of the Tri-Service General Hospital approved this study (IRB: TSGHIRB No. E202416040) on November 4, 2024, and waived the need for individual consent since all identification data were encrypted in the NHIRD. The study was granted review exemption status, and no artificial intelligence tools were used in data analysis or study preparation. As this was a secondary data

analysis of existing administrative data, no direct patient recruitment or contact was performed. The database provides a representative, real-world population sample, allowing for longitudinal follow-up and comprehensive capture of medical events without selection bias inherent in traditional recruitment. No generative artificial intelligence tools were used in any part of the study writing process.

Study Design and Sampled Participants

This retrospective matched-cohort study used inpatient data from January 1, 2000, to December 31, 2015. Each patient with SARS was required to have a diagnosis in an inpatient setting with ICD-9-CM codes 480.8 and 480.9. We selected 285 patients with SARS from 2003 and matched them with 2850 controls at a 1:10 ratio. Additionally, we identified 699 relatives of patients with SARS and matched them with 6990 controls, also at a 1:10 ratio. Propensity score matching was used using variables including gender, age, insurance premium, comorbidities, location, level of care, and index date. Exclusion criteria for the cohorts included individuals with unknown gender and those diagnosed with psychiatric disorders or pneumonia and influenza (ICD-9-CM codes 480-488) before the index date. The index date was defined as the time when individuals were first diagnosed with SARS within the 1-year study period (Table S1 in Multimedia Appendix 1).

All SARS participants and their matched controls were followed from the index date until the occurrence of an injury, including unintentional injuries (traffic accidents, poisoning, medical-related incidents, falls, burns and fires, drowning, suffocation, crushing injuries, adverse drug reactions, and other unintentional injuries), intentional injuries (suicide, or homicide or abuse), or until death, withdrawal from the NHI program, or the end of 2015.

The covariates include sociodemographic characteristics and comorbidities. Sociodemographic characteristics included gender, age (18-44, 45-64, \geq 65 y), education (<12 years; \geq 12 y), monthly insured premiums, urbanization levels, regions of residence, and levels of medical care. The monthly insured premiums have been divided into three categories in New Taiwan Dollars: <18,000, 18,000-34,999, and \geq 35,000. The urbanization level was defined by population and certain indicators of the city's level of development. Level 1

urbanization was defined as having a population greater than 1,250,000 people. Level 2 urbanization was defined as having a population between 500,000 and 1,250,000. Urbanization levels 3 and 4 were defined as having a population between 150,000 and 500,000 and less than 150,000, respectively [43]. The Charlson Comorbidity Index (CCI) is one of the most widely used comorbidity indices [44,45], which consists of 22 conditions [46]. We used the CCI to quantify the comorbidities since it could predict the in-hospital mortality or outcome in patients with severe adult respiratory infection and other infections [47-49].

Statistical Analysis

Statistical analyses were conducted using SPSS (version 22; IBM Corp). Categorical variables were examined with the chi-square test, while continuous variables, expressed as means (SDs), were compared using 2-sample t tests. To evaluate the risk of injuries among patients with and without exposure to patients with SARS or their relatives, competing risk analysis was performed using Fine and Gray's model to calculate adjusted hazard ratios (AHRs) and 95% CIs, adjusting for sociodemographic characteristics and comorbidities. Additionally, Cox proportional hazards regression and sensitivity analyses were used to assess injury risk through the end of 2015. The Kaplan-Meier method was used to compare the cumulative incidence of injuries between the exposed groups and control cohorts, with differences evaluated using the log-rank test. A P value of <.05 was considered statistically significant.

Results

Study Cohort Characteristics

Table 1 presents the demographic and clinical characteristics—including gender, age, education level, monthly insured premiums, urbanization level, region of residence, comorbidities, and levels of medical care—of patients with SARS and their relatives, both with and without SARS. Compared to the control group, there were no significant differences in these covariates between patients with SARS, their relatives, and the controls. The participant flowchart is presented in Figure 1.

Table 1. Baseline demographic and clinical characteristics of survivors of SARS, first-degree relatives, and matched controls in a nationwide retrospective cohort study (Taiwan, 2000-2015)^a.

	Patients with SA	ARS			Relatives of pa	tients with SAR	S	
	Total (n=3135)	With (n=285)	Without (n=2850)	P value ^b	Total (n=7689)	With (n=699)	Without (n=6990)	P value ^b
Gender, n (%)				≥.99				≥.99
Male	1166 (37.19)	106 (37.19)	1060 (37.19)		4422 (57.51)	402 (57.51)	4020 (57.51)	
Female	1969 (62.81)	179 (62.81)	1790 (62.81)		3267 (42.49)	297 (42.49)	2970 (42.49)	
Age (years), mean (SD)	49.67 (12.24)	49.12 (12.18)	49.73 (12.25)	.42	46.72 (12.20)	46.69 (12.15)	46.72 (12.21)	.95
Age groups (years), n	1 (%)			≥.99				≥.99

	Patients with SA	ARS			Relatives of pa	atients with SARS	S	
	Total (n=3135)	With (n=285)	Without (n=2850)	P value ^b	Total (n=7689)	With (n=699)	Without (n=6990)	P value
18-44	1562 (49.82)	142 (49.82)	1420 (49.82)		4411 (57.37)	401 (57.37)	4010 (57.37)	
45-64	1012 (32.28)	92 (32.28)	920 (32.28)		2695 (35.05)	245 (35.05)	2450 (35.05)	
≥65	561 (17.89)	51 (17.89)	510 (17.89)		583 (7.58)	53 (7.58)	530 (7.58)	
Insured premium (New	w Taiwan \$) ^c , n (%	(b)		.48				.98
<18,000	3068 (97.86)	281 (98.60)	2787 (97.79)		7573 (98.49)	689 (98.57)	6884 (98.48)	
18,000-34,999	48 (1.53)	2 (0.70)	46 (1.61)		59 (0.77)	5 (0.72)	54 (0.77)	
≥35,000	19 (0.61)	2 (0.70)	17 (0.60)		57 (0.74)	5 (0.72)	52 (0.74)	
Marital status, n (%)				.38				.92
Without	1465 (46.73)	126 (44.21)	1339 (46.98)		3556 (46.25)	322 (46.07)	3234 (46.27)	
With	1670 (53.27)	159 (55.79)	1511 (53.02)		4133 (53.75)	377 (53.93)	3756 (53.73)	
Education levels (year	rs), n (%)			.95				≥.99
<12	1625 (51.83)	147 (51.58)	1478 (51.86)		3941 (51.26)	359 (51.36)	3582 (51.24)	
≥12	1510 (48.17)	138 (48.42)	1372 (48.14)		3748 (48.74)	340 (48.64)	3408 (48.76)	
CCI_R ^d , n (%)				≥.99				≥.99
0	1738 (55.44)	158 (55.44)	1580 (55.44)		5288 (68.77)	482 (68.96)	4806 (68.76)	
1	638 (20.35)	58 (20.35)	580 (20.35)		1471 (19.13)	131 (18.74)	1340 (19.17)	
≥2	759 (24.21)	69 (24.21)	690 (24.21)		930 (12.10)	86 (12.30)	844 (12.07)	
Location, n (%)				.93				≥.99
Northern Taiwan	1437 (45.84)	132 (46.32)	305 (45.79)		3579 (46.55)	325 (46.49)	3254 (46.55)	
Middle Taiwan	597 (19.04)	54 (18.95)	543 (19.05)		1464 (19.04)	133 (19.03)	1331 (19.04)	
Southern Taiwan	961 (30.65)	87 (30.53)	874 (30.67)		2352 (30.59)	214 (30.62)	2138 (30.59)	
Eastern Taiwan	135 (4.31)	11 (3.86)	124 (4.35)		284 (3.69)	26 (3.72)	258 (3.69)	
Outlying islands	5 (0.16)	1 (0.35)	4 (0.14)		10 (0.13)	1 (0.14)	9 (0.13)	
Urbanization level, n	(%)			.95				≥.99
1 (the highest)	1651 (52.66)	154 (54.04)	1497 (52.53)		4154 (54.03)	378 (54.08)	3776 (54.02)	
2	779 (24.85)	69 (24.21)	710 (24.91)		1875 (24.39)	170 (24.32)	1705 (24.39)	
3	535 (17.07)	46 (16.14)	489 (17.16)		1245 (16.19)	113 (16.17)	1132 (16.19)	
4 (the lowest)	170 (5.42)	16 (5.61)	154 (5.40)		415 (5.40)	38 (5.44)	377 (5.39)	
Level of care, n (%)				.98				≥.99
Hospital center	1634 (52.12)	149 (52.28)	1485 (52.11)		4013 (52.19)	365 (52.22)	3648 (52.19)	
Regional hospital	1170 (37.32)	107 (37.54)	1063 (37.30)		2889 (37.57)	262 (37.48)	2627 (37.58)	
Local hospital	331 (10.56)	29 (10.18)	302 (10.60)		787 (10.24)	72 (10.30)	715 (10.23)	

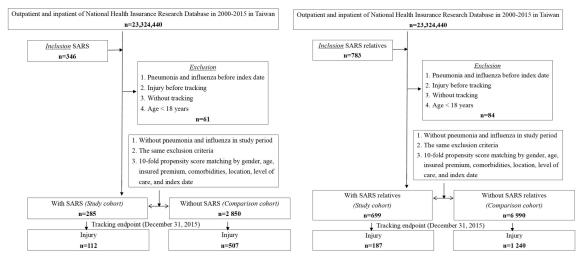
^aThis table presents the demographic and clinical baseline characteristics of survivors of SARS (n=285), their first-degree relatives (n=699), and matched non-SARS controls (n=2850 and n=6990, respectively), extracted from the National Health Insurance Research Database in Taiwan. The data spans from 2000 to 2015 and includes variables such as gender, age, marital status, education level, comorbidity index, insurance premiums, and health care access. The groups were matched 1:10 based on age, gender, comorbidities, and other demographic factors. Statistical significance was assessed using 2-tailed chi-square or Fisher exact tests for categorical variables and 2-tailed *t* tests for continuous variables.

 $^{{}^{\}mathrm{b}}P$ values: chi-square and Fisher exact test on category variables and t test on continue variables.

^cNT\$1=US \$0.035.

 $^{{}^}d\!CCI\!: Charlson\ Comorbidity\ Index, Revised.$

Figure 1. Flowchart detailing the selection of survivors of severe acute respiratory syndrome (SARS) and their first-degree relatives and matched control cohorts from the National Health Insurance Research Database in a nationwide retrospective cohort study conducted in Taiwan (2000-2015).

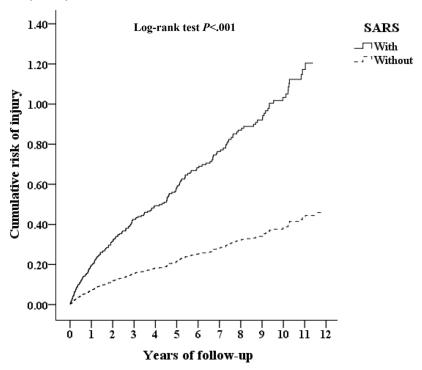


Patients With SARS Kaplan-Meier Curves for the Cumulative Incidence of Injury

In total, 285 patients were diagnosed with SARS during the study period. During the follow-up period, 112 in the

SARS group (n=285) and 507 in the control group (n=2850) developed injury (5560.38 vs 2284.63 per 100,000 personyears). Figure 2 reveals that the difference between the two cohorts in the injury was significant (log-rank test; *P*<.001).

Figure 2. Kaplan-Meier survival curves illustrating the cumulative incidence of injury among the survivors of severe acute respiratory syndrome (SARS) and matched controls aged 18 years or older in a nationwide retrospective cohort study (Taiwan, 2000–2015). Statistical significance was determined using the log-rank test (*P*<.001).

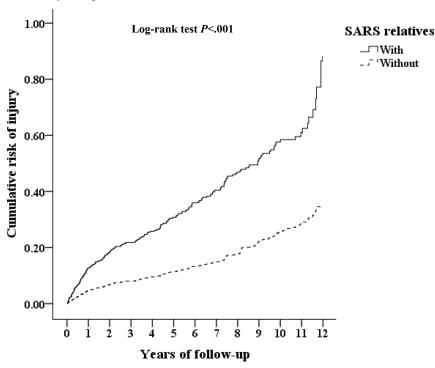


Relatives of Patients With SARS Kaplan-Meier Curves for the Cumulative Incidence of Injury

In total, 699 were relatives of patients with SARS during the study period. During the follow-up period, 187 in the

relatives of patients with SARS group (n=699) and 1240 in the control group (n=6990) developed injury (3044.57 vs 2236.71 per 100,000 person-years). Figure 3 reveals that the difference between the two cohorts in the injury was significant (log-rank test; P<.001).

Figure 3. Kaplan-Meier survival curves demonstrating the cumulative incidence of injury among first-degree relatives of survivors of severe acute respiratory syndrome (SARS) compared to matched controls aged 18 years or older in a nationwide retrospective cohort study (Taiwan, 2000-2015). Statistical significance was evaluated by the log-rank test (*P*<.001).



Years From Patients With SARS and Their Relatives to Injuries

The mean time from the index date to the diagnosis of injuries after diagnosis was 3.60 (SD 3.45) years. The mean years to developed injuries in patients with SARS were 3.47 (SD 3.02) years, which was earlier than in patients without SARS (3.61, SD 3.49 years; Table S2 in Multimedia Appendix 1). The mean time from the index date to the diagnosis of injuries after diagnosis was 3.61 (SD 3.45) years. The mean years to the developed injuries in relatives of patients with SARS were 3.54 (SD 3.07) years, which was earlier than relatives of patients without SARS (3.62, SD 3.49 years; Table S2 in Multimedia Appendix 1).

Analysis of Injury in Patients With SARS and Their Relatives

Table 2 shows the factors of SARS and relatives of patients with SARS, using Fine and Gray's survival analysis, of the factors associated with the risk of injury. In patients with SARS, the crude AHR was 1.684 (95% CI 1.197-2.001; P<.001), and after adjusting for gender, age, education, monthly insured premiums, urbanization levels, regions of residence, comorbidities, and levels of medical care, the adjusted AHR was 1.631 (95% CI 1.184-2.011; P<.001) for injury. In relatives of patients with SARS, the crude AHR was 1.670 (95% CI 1.283-2.025; P<.001), and after adjusting for gender, age, education, monthly insured premiums, urbanization levels, regions of residence, comorbidities, and levels of medical care, the adjusted AHR was 1.572 (95% CI 1.148-1.927; P<.001) for injury. Factors associated with an increased risk of injury among patients with SARS and their relatives included being male, aged 45-64 years or older than 65 years, having a CCI score of ≥2, residing in urbanization levels 1 and 2, and receiving care from medical centers and regional hospitals.

Table 2. AHRs^a for factors associated with injury risk among survivors of severe acute respiratory syndrome (SARS) and their first-degree relatives compared to matched controls using Fine and Gray's competing risk model in a nationwide retrospective cohort study (Taiwan, 2000-2015)^b.

Patients with SARS	Patients v	Patients with SARS					Relatives	Relatives of patients with SARS	ARS			
								La Language Command				
	Competin	Competing risk in the model					Competir	Competing risk in the model	=			
	Crude	95% CI	Ь	Adjusted HR 95% CI	95% CI	Ь	Crude	95% CI	P	Adjusted HR 95% CI	95% CI	P value
	HR^c		value			value	H		value			
SARS (reference: without SARS)	1.684	1.197-2.001	<.001	1.631	1.184-2.011	<.001	1.670	1.283-2.025	<.001 1.572	1.572	1.148-1.927	<.001
Male (reference: female)	1.454	1.396-1.572	<.001	1.386	1.267-1.498	<.001	1.428	1.371-1.555	<.001 1.352	1.352	1.211-1.409	<.001
Age (reference: 18-44 years)												
45-64 years	1.428	1.199-1.426	<.001	1.401	1.184-1.382	<.001 1.411	1.411	1.201-1.529	<.001 1.382	1.382	1.10-1.486	<.001
≥65 years	1.677	1.503-1.786	<.001	1.623	1.490-1.777	<.001 1.637	1.637	1.512-1.803	<.001 1.597	1.597	1.482-1.770	<.001
CCI ^d (reference: CCI=0)												
1	1.325	1.234-1.496	<.001	1.312	1.204-1.452	<.001 1.311	1.311	1.201-1.489	<.001 1.297	1.297	1.184-1.446	<.001
23	1.620	1.591-1.701	<.001	1.607	1.558-1.693	<.001	1.568	1.401-1.698	<.001 1.525	1.525	1.372-1.687	<.001
Urbanization level (reference level: 4)												
1 (the highest)	1.397	1.222-1.489	<.001	1.328	1.201-1.412	<.001 1.375	1.375	1.273-1.586	<.001 1.303	1.303	1.229-1.512	<.001
2	1.286	1.137-1.402	<.001	1.246	1.101-1.354	<.001	1.312	1.146-1.503	<.001 1.289	1.289	1.1111-1.487	<.001
Level of care (reference: local hospital)												
Hospital center	1.979	1.733-2.079	<.001	1.928	1.701-2.016	<.001 1.889	1.889	1.684-2.011	<.001 1.813	1.813	1.607-1.978	<.001
Regional hospital	1.633	1.401-1.976	<.001	1.604	1.389-1.921	<.001	1.606	1.375-1.970	<.001 1.597	1.597	1.342-1.784	<.001

^aAHR: adjusted hazard ratio.

and hospital level of care. The analysis adjusts for baseline characteristics such as demographic variables, comorbidity burden, and access to health care. Data were collected from the National Health Insurance Research Database during the period 2000-2015, with statistical significance set at P<05. first-degree relatives (n=699), compared to matched non-SARS controls. Hazard ratios and 95% CIs are presented for each factor, including SARS exposure, gender, age, comorbidity index, urbanization level, bThis table reports the results of multivariable Cox regression analysis using Fine and Gray's competing risk model to evaluate the factors influencing injury risk among survivors of SARS (n=285) and their

^cHR: hazard ratio.

¹CCI: Charlson Comorbidity Index.

Subgroup Analysis of Unintentional Injuries and Intentional Injuries in the SARS Cohort and Controls

Table S3 in Multimedia Appendix 1 shows that the SARS cohort had an overall AHR of 1.631 (95% CI 1.184-2.011; P<.001). The AHRs for these unintentional injuries were: overall unintentional injuries 1.711 (95% CI 1.204-2.112; P<.001), poisoning 2.701 (95% CI 1.956-4.084; P<.001), and falls 1.524 (95% CI 1.102-1.878; P=.003). For intentional injuries, the overall AHR was 2.232 (95% CI 1.695-2.879; P<.001), with suicide at 2.685 (95% CI 1.947-3.313; P<.001) and homicide or abuse at 1.846 (95% CI 1.341-2.277; P<.001).

Subgroup Analysis of Unintentional Injuries and Intentional Injuries in the Relatives of Patients With SARS Cohort and Controls

Table S4 in Multimedia Appendix 1 presents the AHR for the relatives of patients with SARS cohort, which exhibited a significantly increased overall injury risk with an AHR of 1.572 (95% CI 1.148-1.927; P<.001). In the category of unintentional injuries, the AHRs were notably elevated: overall unintentional injuries had an AHR of 1.742 (95% CI 1.270-2.135; P<.001); traffic-related injuries, 2.003 (95% CI 1.462-2.459; P<.001); poisoning, 1.531 (95% CI 1.120-1.886; P<.001); falls, 1.802 (95% CI 1.324-2.214; P<.001); and crushing injuries, 2.469 (95% CI 1.803-3.026; P<.001). Similarly, intentional injuries demonstrated significantly higher AHR: overall intentional injuries, 2.876 (95% CI 2.101-3.529; P<.001); suicide, 2.197 (95% CI 1.603-2.678; P<.001); and homicide or abuse, 4.163 (95% CI 3.032-5.010; P<.001).

Sensitivity Test for Analysis of the Risk of Injury in Patients With SARS and Their Relatives

Patients with SARS and their relatives were associated with an increased risk of injury compared to the control group. For patients with SARS, the AHR for these injuries was 1.631 (95% CI 1.184-2.011; P<.001). For relatives of patients with SARS, the AHR was 1.572 (95% CI 1.148-1.927; P<.001). Sensitivity analyses revealed that the associations between injury and both the patients with SARS and their relative cohorts remained significant after excluding individuals diagnosed with injury within the first year, as well as when those diagnosed within the first 5 years were excluded (Table 3).

Table 3. Sensitivity analysis of injury risk excluding early injury events in survivors of severe acute respiratory syndrome (SARS) and their first-degree relatives compared to matched controls using Fine and Gray's competing risk model in a nationwide retrospective cohort study (Taiwan, 2000-2015)^a.

	Patient	Patients with SARS	RS		Patients with SARS					Relatives	of patien	Relatives of patients with SARS	RS					
	With			Withou	Without (reference)		Competin	Competing risk in the model		With			Without	Without (reference)	_	Competin	Competing risk in the model	_
Sensitivit y test	Event	Sensitivit Event PYs ^b y test s	Rate I (per 10 ⁵ s PYs)	Event PYs	PYs	Rate (per 10 ⁵ PYs)	AHR ^c	95% CI	P value	Event PYs		Rate (per 10 ⁵ PYs)	Event PYs s		Rate (per 10 ⁵ PYs)	AHR	AHR 95% CI	P value
Overall	112	2014.25	5560.38	507	Overall 112 2014.25 5560.38 507 22,191.7 2284.63 1.631	2284.63	1.631	1.184-2.011	<.001	187	6142.09	3044.57	1240	55,438.5 7	2236.71	1.572	<.001 187 6142.09 3044.57 1240 55,438.5 2236.71 1.572 1.148-1.927	<.001
In the first year excluded	101	1812.80	101 1812.80 5571.49 483	483	19,972.3 2418.35 1.618 0	2418.35	1.618	1.177-2.000	<.001	<.001 167 5	5527.81	3021.09	1189	5527.81 3021.09 1189 49,894.2 2383.04 1.529 5	2383.04	1.529	1.124-1.906	<.001
In the first 5 years excluded	62	1173.25	5284.47	303	1173.25 5284.47 303 12,945.3 2340.61 1.429 3	2340.61	1.429	1.051-1.867	.001	108	3582.70	3582.70 3014.49 803		32,339.1 2483.06 1.498 8	2483.06	1.498	1.095-1.873	<.001

first 5 years of follow-up. Person-years, event rates per 100,000 person-years, and adjusted hazard ratios with 95% CIs are reported. Data were derived from the National Health Insurance Research Database in Taiwan and analyzed using Fine and Gray's model, adjusting for relevant baseline characteristics. ^aThis table displays results of sensitivity analyses evaluating the robustness of injury risk estimates among survivors of SARS and their relatives by excluding injury events occurring within the first year and

^bPY: person-year. ^cAHR: adjusted hazard ratio.

Discussion

Summary of Main Findings

Our study conducted long-term follow-ups of patients with SARS and their relatives. This longitudinal approach provides a comprehensive understanding of the enduring impacts of SARS on both patients and their families. A systematic review conducted in Singapore observed significant decreases in elective, traumatic, and outpatient orthopedic surgeries by 80%, 47%, and 63%, respectively, since the onset of the pandemic. This review found an overall proportional increase in domestic injuries and polytraumas, which was attributed to patients with minor injuries potentially avoiding treatment to minimize their risk of COVID-19 exposure [50]. Additionally, a study from Michigan reported an overall 45% reduction in orthopedic trauma encounters at a level II trauma center following the implementation of a shelter-inplace order [51]. Other research has similarly demonstrated a trend of decreased pediatric and adult trauma and orthopedic admissions since the beginning of the COVID-19 pandemic. However, none of these studies have specifically examined the incidence of injuries or changes in trauma admissions in the northwest Ohio region [52-58].

In this study, we identified several noteworthy findings regarding injury risks within the SARS cohort and their relatives. Specifically, the SARS cohort exhibited a 1.631fold increased risk of overall injuries compared to the control group. Delving into the causes of these injuries, unintentional injuries presented a 1.711-fold increase, poisoning incidents rose by 2.701-fold, and falls were associated with a 1.524-fold higher risk. Additionally, the cohort of relatives of patients with SARS demonstrated a 1.572-fold elevated risk of overall injuries. Breaking down the injury causes within this group, unintentional injuries showed a 1.742-fold increase, traffic-related injuries doubled with a 2.003-fold rise, poisoning incidents were up by 1.531-fold, medical-related injuries increased by 1.362-fold, falls were 1.802-fold more likely, and crushing injuries saw the highest increase at 2.469-fold. In this study, "relatives" are defined as first-degree family members (spouses, children, parents, or siblings) sharing the same NHI household registration [7]. Because the NHIRD does not capture cohabitation or caregiving roles, we cannot determine whether these relatives lived with or provided direct care to the patients with SARS. Subgroup analysis (Table S4 in Multimedia Appendix 1) shows that relatives of deceased patients with SARS had an AHR of 2.34 (95% CI 2.10-2.61), compared to 1.78 (95% CI 1.56-2.03) for relatives of surviving patients. The higher risk among bereaved relatives may reflect grief-related trauma and family disruption, though increased health care interactions or reporting bias cannot be excluded. Future research should incorporate data on household residency and caregiving responsibilities—such as via linkage to household registries or targeted surveys-to clarify the pathways by which SARS exposure affects relatives' injury risk.

Detailed Discussion and Interpretation

Interestingly, in other studies, we have found that falls encounters increased during the COVID-19 pandemic. The impact of COVID-19 on the older population included not only detrimental health effects but also societal effects, such as increased isolation leading to higher rates of depression, anxiety, and dementia [5]. We hypothesize that the incidence of falls increased among the older population due to the decreased availability of professional caregivers during COVID-19. With fewer family members and professional help available, older patients may have been more susceptible to falling. Additionally, in this study, the risk of falls remained elevated 15 years after tracking patients with SARS, which we attribute to the physical weakness resulting from the illness. In contrast, among relatives of patients with SARS, the risk of falls did not increase. These findings highlight the long-term physical vulnerabilities in patients with SARS and suggest that the broader societal disruptions caused by pandemics can have lasting impacts on injury risks within vulnerable populations. The literature indicates that fatigue was the most common symptom among survivors of SARS 18 years after discharge, with osteoporosis and femoral head necrosis being the primary sequelae. The respiratory and hip function scores of survivors of SARS were significantly lower compared to those of the control group. Physical and social functioning at 18 years postdischarge showed improvement compared to that at 12 years but still remained inferior to the controls. Emotional and mental health were fully recovered. Lung lesions detected by computed tomography scans remained consistent at 18 years, particularly in the right upper lobe and left lower lobe. Plasma multiomics analysis indicated abnormal metabolism of amino acids and lipids, an enhanced host immune defense response to bacteria and external stimuli, B-cell activation, and increased cytotoxicity of CD8+ T cells, while showing impaired antigen presentation capacity of CD4+ T cells [6,59-61]. These factors may contribute to the increased risk of falls among patients with SARS.

The persistence of suicide and homicide, or abuse risk morbidities among survivors of SARS and their relatives who participated in this study was alarming. This study identified a particularly noteworthy finding: among patients with SARS, the risk of homicide or abuse was elevated by a factor of 1.846. However, this risk was significantly higher among the relatives of patients with SARS, increasing to 4.163 times. Reviewing results from previous post-SARS cohorts, psychiatric morbidities measured by standardized questionnaires ranged from 10% to 35% in the acute phase of the infection (from the acute phase to 1 month after SARS) [62,63], increasing to 64% at the 1-year follow-up [64]. In the context of the COVID-19 pandemic, research has demonstrated a significant change in the number of nonaccidental trauma encounters, with more cases occurring during the pre-COVID-19 period than during COVID-19 among both adult (15 vs 11 cases) and pediatric (29 vs 9 cases) populations. There is substantial concern that individuals may have been subjected to increased neglect and abuse due to the heightened time spent at home during COVID-19 [8]. Regarding the observed decrease in nonaccidental trauma encounters during COVID-19, concerns have been raised about whether this reduction was due to fewer patients seeking treatment for these traumas rather than a true decrease in the incidence of such events. During the pandemic, time spent engaging in activities outside the home-where nonaccidental trauma could be identified and reported-considerably decreased. In addition to the psychosocial stressors associated with SARS and COVID-19, the long-term adverse health outcomes experienced by survivors of SARS may also serve as a risk factor for accidental injuries among their relatives [65,66]. The rise in suicide rates following the SARS outbreak was a critical public health concern. Previous studies have explored the potential increased risk of suicide among older populations [3,4] and emergency department visitors [67] during and after the SARS outbreak. To our knowledge, this study represents the first investigation into suicide specifically among survivors of SARS. As previously discussed, the association between SARS and suicide risk was found to be statistically significant, even after excluding suicides occurring within the first year postinfection. However, this association was not significant after excluding suicides that occurred within the first 5 years. These findings highlight the importance of vigilant assessment of suicide risk in the years following SARS infection.

To elucidate pathways linking SARS exposure to elevated injury risks, we propose a 3-fold framework. Biological: chronic fatigue, pulmonary fibrosis, and immune dysregulation in survivors of SARS may impair physical resilience and coordination, contributing to a higher fall risk (AHR 1.524, 95% CI 1.102-1.878; Table S3 in Multimedia Appendix 1) and paralleling neurological sequela observed in survivors of COVID-19 [68]. Psychological: quarantine stress, fear of morbidity or mortality, and economic hardship exacerbate mental-health complications, elevating self-harm risk (suicide AHR 2.685, 95% CI 1.947-3.313; Table S3 in Multimedia Appendix 1) and accidental injuries. Social: reduced social support and disrupted routines further heighten vulnerability, particularly among isolated individuals. Preventive interventions: regular physical activity has demonstrated efficacy in preventing postinfection mental-health complications [69], suggesting a modifiable strategy to mitigate injury risk in both SARS and COVID-19 populations [70,71].

Factors contributing to this increased burden may include stress from quarantine and isolation, fear of uncertainty and death, impaired physical health following severe viral infection, the economic consequences of SARS, and even adverse emotional responses to SARS-related information [71-73]. These factors collectively underscore the necessity for careful monitoring and proactive intervention to mitigate psychiatric risks in affected populations.

Limitations

First, although we achieved an excellent balance on observed covariates—gender, age, insurance premium, CCI, geographic location, level of care, and index date (all *P*>.05; Table 1)—our retrospective matched-cohort design cannot

definitively establish causality. Unmeasured confounders—such as occupational exposures (eg, health care workers at hospitals and airport or transit staffs) or underlying socioeconomic and environmental factors—may bias our estimates. Indeed, the AHR for homicide or abuse among relatives (AHR 4.163; 95% CI 3.032-5.010) substantially exceeds that for patients (AHR 1.846; 95% CI 1.341-2.277; Table S4 in Multimedia Appendix 1), highlighting potential residual confounding. Moreover, our analysis is based exclusively on Taiwan's 2003 SARS outbreak, which featured distinctive health care responses, social-distancing policies, and cultural norms; therefore, generalizing to other populations or to the COVID-19 pandemic—despite emerging parallels in postinfection neurological sequela—requires caution [68].

Second, the NHIRD does not capture validated injury severity measures (eg, standardized scales), nor does it include laboratory or imaging results or confirmatory diagnostic-test data, limiting our capacity to explore underlying injury mechanisms or to stratify outcomes by clinical severity [40].

Third, patients treated for SARS may have been more health-conscious, potentially leading to an overestimation of injury incidence. Additionally, clinicians may have exercised heightened vigilance in managing survivors of SARS, possibly increasing inpatient referrals. However, sensitivity analyses excluding injuries occurring within the first year and first 5 years after SARS still showed significantly elevated risks (AHR 1.618, 95% CI 1.177-2.000; AHR 1.429, 95% CI 1.051-1.867). This suggests that differential treatment decisions alone are unlikely to explain the higher hazard ratios observed.

Finally, our 15-year follow-up spanned periods of major shifts in health care delivery (eg, rapid adoption of telemedicine and reorganization of hospital services for infection control), social-support structures (eg, community quarantine measures and establishment of outreach networks), and injury-reporting workflows (eg, revised triage protocols and updated insurance-claim coding). Such changes likely affected both care-seeking behaviors and the administrative capture of injuries; for instance, outpatient visits decreased by approximately 45% during the SARS outbreak [74], indicating potential underascertainment of minor injuries. Similar patterns of health care avoidance have been observed during the COVID-19 pandemic, highlighting the importance of contextualizing our results within evolving health care and reporting environments.

Future research should include more detailed clinical data, address potential confounders, and consider expanding the cohort to include family members and health care workers for a more comprehensive understanding.

Conclusions

This study found that survivors of SARS and their relatives had a significantly increased risk of both unintentional and intentional injuries, persisting over 15 years. Relatives faced a heightened risk of homicide or abuse, while survivors showed increased risks of poisoning, falls, and suicide. These findings

highlight the need for clinicians to closely monitor the psychological and injury risks in survivors of SARS and their to ensure adequate mental health and safety support.

Acknowledgments

The authors would like to express their gratitude and appreciation to the Health and Welfare Data Science Center, Ministry of Health and Welfare, Taiwan, for providing access to the National Health Insurance Research Database. This research was funded by the Tri-Service General Hospital (grants TSGH-B-114022 and TSGH-A-114010 special plan).

Data Availability

This study uses third-party data. Taiwan launched a single-payer National Health Insurance (NHI) program on March 1, 1995. The database of this program contains registration files and original claim data for reimbursement. Large computerized databases are derived from this system by the NHI Administration. Investigators interested may submit a formal proposal to the NHI Research Database. The authors confirm they did not have any special access privileges.

Authors' Contributions

CS wrote the Introduction, Discussion, and Conclusion sections, contributed to drafting the Abstract, and prepared Table 1 and Table 2. CHC implemented the software, verified data accuracy, and contributed to data validation. FHL participated in methodology development, conducted data validation, and supported the interpretation of statistical results. DYN cowrote the Results section and contributed to drafting the Abstract and Supplementary Tables S2 to S4. CAS designed the study methodology, curated the dataset, and constructed Table S1. CHT contributed to the study design and performed cohort matching and data curation. LYF conceptualized the study, supervised the research process, and critically reviewed the manuscript. WCC coconceptualized the study, coordinated the research project, secured funding, and reviewed and edited the final manuscript. All authors reviewed and approved the final version of the manuscript

Conflicts of Interest

None declared.

Multimedia Appendix 1

Supplementary tables.

[DOCX File (Microsoft Word File), 28 KB-Multimedia Appendix 1]

References

- Ruuskanen O, Lahti E, Jennings LC, Murdoch DR. Viral pneumonia. Lancet. Apr 9, 2011;377(9773):1264-1275. [doi: 10.1016/S0140-6736(10)61459-6] [Medline: 21435708]
- 2. Acute respiratory syndrome. China, Hong Kong Special Administrative Region of China, and Viet Nam. Wkly Epidemiol Rec. Mar 14, 2003;78(11):73-74. [Medline: 12674023]
- 3. Chan SMS, Chiu FKH, Lam CWL, Leung PYV, Conwell Y. Elderly suicide and the 2003 SARS epidemic in Hong Kong. Int J Geriatr Psychiatry. Feb 2006;21(2):113-118. [doi: 10.1002/gps.1432] [Medline: 16416469]
- 4. Cheung YT, Chau PH, Yip PSF. A revisit on older adults suicides and severe acute respiratory syndrome (SARS) epidemic in Hong Kong. Int J Geriatr Psychiatry. Dec 2008;23(12):1231-1238. [doi: 10.1002/gps.2056] [Medline: 18500689]
- 5. Abbasi J. Social isolation—the other COVID-19 threat in nursing homes. JAMA. Aug 18, 2020;324(7):619. [doi: <u>10.1001/jama.2020.13484</u>]
- 6. Dhawan RT, Gopalan D, Howard L, et al. Beyond the clot: perfusion imaging of the pulmonary vasculature after COVID-19. Lancet Respir Med. Jan 2021;9(1):107-116. [doi: 10.1016/S2213-2600(20)30407-0] [Medline: 33217366]
- 7. Tzeng NS, Chung CH, Chang CC, et al. What could we learn from SARS when facing the mental health issues related to the COVID-19 outbreak? A nationwide cohort study in Taiwan. Transl Psychiatry. Oct 6, 2020;10(1):339. [doi: 10.1038/s41398-020-01021-y] [Medline: 33024072]
- 8. Sanford EL, Zagory J, Blackwell JM, Szmuk P, Ryan M, Ambardekar A. Changes in pediatric trauma during COVID-19 stay-at-home epoch at a tertiary pediatric hospital. J Pediatr Surg. May 2021;56(5):918-922. [doi: 10.1016/j.jpedsurg. 2021.01.020] [Medline: 33516579]
- 9. Panagea E, Messinis L, Petri MC, et al. Neurocognitive Impairment in long COVID: a systematic review. Arch Clin Neuropsychol. Jan 21, 2025;40(1):125-149. [doi: 10.1093/arclin/acae042] [Medline: 38850628]
- 10. Haunhorst S, Bloch W, Wagner H, et al. Long COVID: a narrative review of the clinical aftermaths of COVID-19 with a focus on the putative pathophysiology and aspects of physical activity. Oxf Open Immunol. 2022;3(1):iqac006. [doi: 10.1093/oxfimm/iqac006] [Medline: 36846561]
- 11. Subramaniam A, Shekar K, Afroz A, et al. Frailty and mortality associations in patients with COVID-19: a systematic review and meta-analysis. Intern Med J. May 2022;52(5):724-739. [doi: 10.1111/imj.15698] [Medline: 35066970]

- 12. Ceban F, Ling S, Lui LMW, et al. Fatigue and cognitive impairment in post-COVID-19 syndrome: a systematic review and meta-analysis. Brain Behav Immun. Mar 2022;101:93-135. [doi: 10.1016/j.bbi.2021.12.020] [Medline: 34973396]
- 13. Taquet M, Geddes JR, Husain M, Luciano S, Harrison PJ. 6-month neurological and psychiatric outcomes in 236 379 survivors of COVID-19: a retrospective cohort study using electronic health records. Lancet Psychiatry. May 2021;8(5):416-427. [doi: 10.1016/S2215-0366(21)00084-5] [Medline: 33836148]
- 14. AlRasheed MM, Fekih-Romdhane F, Jahrami H, et al. The prevalence and severity of insomnia symptoms during COVID-19: a global systematic review and individual participant data meta-analysis. Sleep Med. Dec 2022;100:7-23. [doi: 10.1016/j.sleep.2022.06.020] [Medline: 36030616]
- 15. Arenas DJ, Thomas A, Wang J, DeLisser HM. A systematic review and meta-analysis of depression, anxiety, and sleep disorders in US adults with food insecurity. J Gen Intern Med. Dec 2019;34(12):2874-2882. [doi: 10.1007/s11606-019-05202-4] [Medline: 31385212]
- 16. Paterson RW, Brown RL, Benjamin L, et al. The emerging spectrum of COVID-19 neurology: clinical, radiological and laboratory findings. Brain (Bacau). Oct 1, 2020;143(10):3104-3120. [doi: 10.1093/brain/awaa240]
- 17. Ambrose AF, Paul G, Hausdorff JM. Risk factors for falls among older adults: a review of the literature. Maturitas. May 2013;75(1):51-61. [doi: 10.1016/j.maturitas.2013.02.009] [Medline: 23523272]
- 18. Woodrow M, Carey C, Ziauddeen N, et al. Systematic review of the prevalence of long COVID. Open Forum Infect Dis. Jul 2023;10(7):ofad233. [doi: 10.1093/ofid/ofad233] [Medline: 37404951]
- 19. Seppala LJ, Wermelink A, de Vries M, et al. Fall-risk-increasing drugs: a systematic review and meta-analysis: ii. psychotropics. J Am Med Dir Assoc. Apr 2018;19(4):371. [doi: 10.1016/j.jamda.2017.12.098] [Medline: 29402652]
- 20. Keshta AS, Mallah SI, Al Zubaidi K, et al. COVID-19 versus SARS: a comparative review. J Infect Public Health. Jul 2021;14(7):967-977. [doi: 10.1016/j.jiph.2021.04.007] [Medline: 34130121]
- 21. Low ZY, Yip AJW, Sharma A, Lal SK. SARS coronavirus outbreaks past and present-a comparative analysis of SARS-CoV-2 and its predecessors. Virus Genes. Aug 2021;57(4):307-317. [doi: 10.1007/s11262-021-01846-9] [Medline: 34061288]
- 22. Chan WSH. Taiwan's healthcare report 2010. EPMA J. Dec 2010;1(4):563-585. [doi: 10.1007/s13167-010-0056-8] [Medline: 23199110]
- 23. Chien WC, Chung CH, Lin FH, Chang HA, Kao YC, Tzeng NS. Is weight control surgery associated with increased risk of newly onset psychiatric disorders? A population-based, matched cohort study in Taiwan. J Med Sci. 2017;37(4):137. [doi: 10.4103/jmedsci.jmedsci 94 16]
- 24. Chen TY, Huang CH, Chung CH, et al. Sex and age differences in the association between anxiety disorders and narcolepsy: a nationwide population-based case control study. J Affect Disord. Mar 1, 2020;264:130-137. [doi: 10.1016/j.jad.2019.12.010] [Medline: 32056742]
- 25. Lin CH, Chien WC, Chung CH, et al. Increased risk of dementia in patients with genital warts: a nationwide cohort study in Taiwan. J Dermatol. May 2020;47(5):503-511. [doi: 10.1111/1346-8138.15277] [Medline: 32189395]
- 26. Lin YC, Chen TY, Chien WC, et al. Stimulants associated with reduced risk of hospitalization for motor vehicle accident injury in patients with obstructive sleep apnea—a nationwide cohort study. BMC Pulm Med. Feb 3, 2020;20(1):28. [doi: 10.1186/s12890-019-1041-1] [Medline: 32013932]
- 27. Liu YP, Chien WC, Chung CH, Chang HA, Kao YC, Tzeng NS. Are anticholinergic medications associated with increased risk of dementia and behavioral and psychological symptoms of dementia? a nationwide 15-year follow-up cohort study in Taiwan. Front Pharmacol. 2020;11:30. [doi: 10.3389/fphar.2020.00030] [Medline: 32116707]
- 28. Wan FJ, Chien WC, Chung CH, Yang YJ, Tzeng NS. Association between traumatic spinal cord injury and affective and other psychiatric disorders—a nationwide cohort study and effects of rehabilitation therapies. J Affect Disord. Mar 15, 2020;265:381-388. [doi: 10.1016/j.jad.2020.01.063] [Medline: 32090763]
- 29. Wang DS, Chung CH, Chang HA, et al. Association between child abuse exposure and the risk of psychiatric disorders: a nationwide cohort study in Taiwan. Child Abuse Negl. Mar 2020;101:104362. [doi: 10.1016/j.chiabu.2020.104362]
- 30. Yang CC, Chien WC, Chung CH, et al. No association between human immunodeficiency virus infections and dementia: a nationwide cohort study In Taiwan. Neuropsychiatr Dis Treat. 2019;15:3155-3166. [doi: 10.2147/NDT.S225584] [Medline: 31814723]
- 31. Yeh TC, Chien WC, Chung CH, et al. Psychiatric disorders after traumatic brain injury: a nationwide population-based cohort study and the effects of rehabilitation therapies. Arch Phys Med Rehabil. May 2020;101(5):822-831. [doi: 10.1016/j.apmr.2019.12.005] [Medline: 31917196]
- 32. Kao LC, Chien WC, Chung CH, et al. The newly diagnosed amnestic disorders and dementia: a nationwide, cohort study in Taiwan. Taiwan J Psychiatry. 2018;32(1):18-28. URL: https://www.sop.org.tw/sop_journal/Upload_files/32_1/02.pdf [Accessed 2025-06-24]

- 33. Chiou YE, Chien WC, Chung CH, et al. New users of herbal medicine containing aristolochic acids and the risk of dementia in the elderly: a nationwide, population-based study in Taiwan. Neuropsychiatr Dis Treat. 2020;16:1493-1504. [doi: 10.2147/NDT.S250659] [Medline: 32606702]
- 34. Chien WC, Chung CH, Lin FH, et al. The risk of injury in adults with attention-deficit hyperactivity disorder: a nationwide, matched-cohort, population-based study in Taiwan. Res Dev Disabil. Jun 2017;65:57-73. [doi: 10.1016/j. ridd.2017.04.011] [Medline: 28458048]
- 35. Chien WC, Chung CH, Lai CH, Chou CH. A retrospective population-based study of injury types among elderly in Taiwan. Int J Inj Contr Saf Promot. 2014;21(1):3-8. [doi: 10.1080/17457300.2012.717084] [Medline: 22924733]
- 36. Chang SY, Chien WC, Chung CH, et al. Risk of dementia after charcoal-burning suicide attempts: a nationwide cohort study in Taiwan. J Investig Med. Oct 2018;66(7):1070-1082. [doi: 10.1136/jim-2018-000759] [Medline: 29884660]
- 37. Chu CW, Chien WC, Chung CH, et al. Electroconvulsive therapy and risk of dementia—a nationwide cohort study in Taiwan. Front Psychiatry. 2018;9:397. [doi: 10.3389/fpsyt.2018.00397] [Medline: 30245639]
- 38. Tzeng NS, Chung CH, Lin FH, et al. Anti-herpetic medications and reduced risk of dementia in patients with herpes simplex virus infections—a nationwide, population-based cohort study in Taiwan. Neurotherapeutics. Apr 2018;15(2):417-429. [doi: 10.1007/s13311-018-0611-x] [Medline: 29488144]
- 39. Tzeng NS, Chung CH, Lin FH, et al. Risk of dementia in adults with ADHD: a nationwide, population-based cohort study in Taiwan. J Atten Disord. Jul 2019;23(9):995-1006. [doi: 10.1177/1087054717714057] [Medline: 28629260]
- 40. Hsieh CY, Su CC, Shao SC, et al. Taiwan's National Health Insurance Research Database: past and future. Clin Epidemiol. 2019;11:349-358. [doi: 10.2147/CLEP.S196293] [Medline: 31118821]
- 41. Cheng CL, Kao YHY, Lin SJ, Lee CH, Lai ML. Validation of the National Health Insurance Research Database with ischemic stroke cases in Taiwan. Pharmacoepidemiol Drug Saf. Mar 2011;20(3):236-242. [doi: 10.1002/pds.2087] [Medline: 21351304]
- 42. Liang JA, Sun LM, Muo CH, Sung FC, Chang SN, Kao CH. The analysis of depression and subsequent cancer risk in Taiwan. Cancer Epidemiol Biomarkers Prev. Mar 1, 2011;20(3):473-475. [doi: 10.1158/1055-9965.EPI-10-1280]
- 43. Chang CY, Chen WL, Liou YF, et al. Increased risk of major depression in the three years following a femoral neck fracture—a national population-based follow-up study. PLoS One. 2014;9(3):e89867. [doi: 10.1371/journal.pone. 0089867] [Medline: 24626193]
- 44. Charlson ME, Pompei P, Ales KL, MacKenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. J Chronic Dis. 1987;40(5):373-383. [doi: 10.1016/0021-9681(87)90171-8] [Medline: 3558716]
- 45. de Groot V, Beckerman H, Lankhorst GJ, Bouter LM. How to measure comorbidity. a critical review of available methods. J Clin Epidemiol. Mar 2003;56(3):221-229. [doi: 10.1016/s0895-4356(02)00585-1] [Medline: 12725876]
- 46. Charlson M, Szatrowski TP, Peterson J, Gold J. Validation of a combined comorbidity index. J Clin Epidemiol. Nov 1994;47(11):1245-1251. [doi: 10.1016/0895-4356(94)90129-5] [Medline: 7722560]
- 47. Ternavasio-de la Vega HG, Castaño-Romero F, Ragozzino S, et al. The updated Charlson Comorbidity Index is a useful predictor of mortality in patients with *Staphylococcus aureus* bacteraemia . Epidemiol Infect. Dec 2018;146(16):2122-2130. [doi: 10.1017/S0950268818002480]
- 48. Baiomi A, Abbas H, Pirzada U, Zaidi B, Saturno D, Nayudu SK. Charlson Comorbidity Index (CCI): an independent predictor of outcomes in Clostridium difficile infection (CDI). Am J Gastroenterol. Oct 2018;113(Supplement):S1525. [doi: 10.14309/00000434-201810001-02742]
- 49. Setter NW, Peres ML, de Almeida BMM, Petterle RR, Raboni SM. Charlson Comorbidity Index scores and in-hospital prognosis of patients with severe acute respiratory infections. Intern Med J. Jun 2020;50(6):691-697. [doi: 10.1111/imj. 14398] [Medline: 31180163]
- 50. Ow ZGW, Cheong CK, Chin YH, Chin BZ. A look at the global impact of SARS CoV-2 on orthopedic services. J Clin Orthop Trauma. Jan 2021;12(1):33-39. [doi: 10.1016/j.jcot.2020.10.052] [Medline: 33191995]
- 51. Stoker S, McDaniel D, Crean T, et al. Effect of shelter-in-place orders and the COVID-19 pandemic on orthopaedic trauma at a community level ii trauma center. J Orthop Trauma. Sep 2020;34(9):e336-e342. [doi: 10.1097/BOT. 000000000001860] [Medline: 32815848]
- 52. Bram JT, Johnson MA, Magee LC, et al. Where have all the fractures gone? The epidemiology of pediatric fractures during the COVID-19 pandemic. J Pediatr Orthop. 2020;40(8):373-379. [doi: 10.1097/BPO.0000000000001600]
- 53. Druel T, Andeol Q, Rongieras F, Bertani A, Bordes M, Alvernhe A. Evaluation of containment measures' effect on orthopaedic trauma surgery during the COVID-19 pandemic: a retrospective comparison between 2019 and 2020. Int Orthop. Nov 2020;44(11):2229-2234. [doi: 10.1007/s00264-020-04712-9] [Medline: 32696333]

- 54. Kale NN, Patel AH, Leddy MJ 3rd, Savoie FH 3rd, Sherman WF. The effect of COVID-19 on orthopedic practices and surgeons in Louisiana. Orthopedics. Nov 1, 2020;43(6):351-355. [doi: 10.3928/01477447-20201023-01] [Medline: 33211903]
- 55. Fyllos A, Varitimidis S, Papageorgiou F, et al. Upper extremity emergencies during SARS-COV-2 pandemic: turnout trends. Injury. Mar 2021;52(3):387-394. [doi: 10.1016/j.injury.2020.11.045] [Medline: 33246643]
- 56. Gumina S, Proietti R, Polizzotti G, Carbone S, Candela V. The impact of COVID-19 on shoulder and elbow trauma: an Italian survey. J Shoulder Elbow Surg. Sep 2020;29(9):1737-1742. [doi: 10.1016/j.jse.2020.05.003] [Medline: 32713663]
- 57. Lubbe RJ, Miller J, Roehr CA, et al. Effect of statewide social distancing and stay-at-home directives on orthopaedic trauma at a southwestern level 1 trauma center during the COVID-19 pandemic. J Orthop Trauma. Sep 2020;34(9):e343-e348. [doi: 10.1097/BOT.0000000000001890] [Medline: 32815849]
- 58. Wallace CN, Kontoghiorghe C, Kayani B, Chang JS, Haddad FS. The impact of COVID-19 on trauma and orthopaedic surgery in the United Kingdom. Bone Jt Open. Jul 2020;1(7):420-423. [doi: 10.1302/2633-1462.17.BJO-2020-0102] [Medline: 33215133]
- 59. Li K, Wu Q, Li H, et al. Multiomic characterisation of the long-term sequelae of SARS survivors: a clinical observational study. EClinicalMedicine. Apr 2023;58:101884. [doi: 10.1016/j.eclinm.2023.101884] [Medline: 36873427]
- 60. Tansey CM, Louie M, Loeb M. One-year outcomes and health care utilization in survivors of severe acute respiratory syndrome. Arch Intern Med. Jun 25, 2007;167(12):1312. [doi: 10.1001/archinte.167.12.1312] [Medline: 17592106]
- 61. Lam MHB, Wing YK, Yu MWM, et al. Mental morbidities and chronic fatigue in severe acute respiratory syndrome survivors: long-term follow-up. Arch Intern Med. Dec 14, 2009;169(22):2142-2147. [doi: 10.1001/archinternmed.2009.384] [Medline: 20008700]
- 62. Wu KK, Chan SK, Ma TM. Posttraumatic stress after SARS. Emerg Infect Dis. Aug 2005;11(8):1297-1300. [doi: <u>10.3201/eid1108.041083</u>] [Medline: <u>16102324</u>]
- 63. Sheng B, Cheng SKW, Lau KK, Li HL, Chan ELY. The effects of disease severity, use of corticosteroids and social factors on neuropsychiatric complaints in severe acute respiratory syndrome (SARS) patients at acute and convalescent phases. Eur Psychiatry. May 2005;20(3):236-242. [doi: 10.1016/j.eurpsy.2004.06.023] [Medline: 15935422]
- 64. Lee AM, Wong J, McAlonan GM, et al. Stress and psychological distress among SARS survivors 1 year after the outbreak. Can J Psychiatry. Apr 2007;52(4):233-240. [doi: 10.1177/070674370705200405] [Medline: 17500304]
- 65. Lee DTS, Wing YK, Leung HCM, et al. Factors associated with psychosis among patients with severe acute respiratory syndrome: a case-control study. Clin Infect Dis. Oct 15, 2004;39(8):1247-1249. [doi: 10.1086/424016] [Medline: 15486852]
- 66. Phua DH, Tang HK, Tham KY. Coping responses of emergency physicians and nurses to the 2003 severe acute respiratory syndrome outbreak. Acad Emerg Med. Apr 2005;12(4):322-328. [doi: 10.1197/j.aem.2004.11.015] [Medline: 15805323]
- 67. Huang CC, Yen DHT, Huang HH, et al. Impact of severe acute respiratory syndrome (SARS) outbreaks on the use of emergency department medical resources. J Chin Med Assoc. Jun 2005;68(6):254-259. [doi: 10.1016/S1726-4901(09)70146-7] [Medline: 15984818]
- 68. Park JM, Woo W, Lee SC, et al. Prevalence and mortality risk of neurological disorders during the COVID-19 pandemic: an umbrella review of the current evidence. Neuroepidemiology. 2023;57(3):129-147. [doi: 10.1159/000530536] [Medline: 37044073]
- 69. Rahmati M, Lee S, Yon DK, et al. Physical activity and prevention of mental health complications: an umbrella review. Neurosci Biobehav Rev. May 2024;160:105641. [doi: 10.1016/j.neubiorev.2024.105641] [Medline: 38527637]
- 70. Chua SE, Cheung V, McAlonan GM, et al. Stress and psychological impact on SARS patients during the outbreak. Can J Psychiatry. Jun 2004;49(6):385-390. [doi: 10.1177/070674370404900607] [Medline: 15283533]
- 71. Tsang HWH, Scudds RJ, Chan EYL. Psychosocial impact of SARS. Emerg Infect Dis. Jul 2004;10(7):1326-1327. [doi: 10.3201/eid1007.040090] [Medline: 15338536]
- 72. Sim K, Chua HC. The psychological impact of SARS: a matter of heart and mind. CMAJ. Mar 2, 2004;170(5):811-812. [doi: 10.1503/cmaj.1032003] [Medline: 14993176]
- 73. Sin SS, Huak CY. Psychological impact of the SARS outbreak on a Singaporean rehabilitation department. Int J Ther Rehabil. Sep 2004;11(9):417-424. [doi: 10.12968/ijtr.2004.11.9.19589]
- 74. Chu D, Chen RC, Ku CY, Chou P. The impact of SARS on hospital performance. BMC Health Serv Res. Nov 6, 2008;8(1):228. [doi: 10.1186/1472-6963-8-228] [Medline: 18990210]

Abbreviations

AHR: adjusted hazard ratio

CCI: Charlson Comorbidity Index

ICD-9-CM: International Classification of Diseases, Ninth Revision, Clinical Modification

NHI: National Health Insurance

NHIRD: National Health Insurance Research Database

SARS: severe acute respiratory syndrome

Edited by Amaryllis Mavragani, Travis Sanchez; peer-reviewed by Seung Won Lee, Wei Yang; submitted 26.12.2024; final revised version received 05.06.2025; accepted 08.06.2025; published 08.07.2025

<u>Please cite as:</u>

Sung C, Chung CH, Sun CA, Tsao CH, Ng DY, Weng TH, Fann LY, Lin FH, Chien WC

COVID-19-Relevant Insights Into the Elevated Risk of Accidental Injuries in Survivors of SARS and Their Relatives in

Taiwan: Retrospective Cohort Study

JMIR Public Health Surveill 2025;11:e70608 URL: https://publichealth.jmir.org/2025/1/e70608

doi: 10.2196/70608

© Chieh Sung, Chi-Hsiang Chung, Chien-An Sun, Chang-Huei Tsao, Daphne Yih Ng, Tsu-Hsuan Weng, Li-Yun Fann, Fu-Huang Lin, Wu-Chien Chien. Originally published in JMIR Public Health and Surveillance (https://publichealth.jmir.org), 08.07.2025. This is an open-access article distributed under the terms of the Creative Commons Attribution License (https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work, first published in JMIR Public Health and Surveillance, is properly cited. The complete bibliographic information, a link to the original publication on https://publichealth.jmir.org, as well as this copyright and license information must be included.